Appl. No. 10/801,078 Amdt. dated April 6, 2010 Reply to Restriction of November 15, 2009

Amendments to the Claims:

This listing of claims will replace all prior versions, and listings of claims in the application:

Listing of Claims:

1-15 (canceled)

16. (Currently amended) A method of treating a human subject suffering loss of photoreceptor function, said method comprising:

administering an effective amount of a 9-cis retinal derivative, wherein said derivative is not 9-cis retinal and is a modification of the polyene chain but with retention of the polyene chain length and retention of the 9-cis bond, in a pharmaceutically acceptable vehicle to a human subject, with autosomal dominant retinitis pigmentosa due to expression of a mutant opsin protein with a substitution of Proline 23 by Histidine (P23H mutant opsin protein), to treat loss of photoreceptor function in said subject,

wherein 9-cis retinal has the structure of

- 17. (Previously presented) The method of claim 16, wherein the retinal derivative is orally administered to the human subject.
- 18. (Previously presented) The method of claim 16, wherein the retinal derivative is locally administered to the human subject.

19-34. (canceled)

- 35. (Previously presented) The method of claim 18, wherein the retinal derivative is locally administered by eye drops.
- 36. (Previously presented) The method of claim 18, wherein the retinal derivative is locally administered by intraocular injection.
- 37. (Previously presented) The method of claim 18, wherein the retinal derivative is locally administered by periocular injection.

38-48. (canceled)

- 49. (Previously presented) The method of claim 16, wherein the subject endogenously forms rhodopsin, from opsin and endogenous 11-cis-retinal, as a visual pigment.
- 50. (Previously presented) The method of claim 18, wherein the subject endogenously forms rhodopsin, from opsin and endogenous 11-cis-retinal, as a visual pigment.
- 51. (Previously presented) The method of claim 16, further comprising identifying the subject as expressing a mutant opsin protein with a substitution of Proline 23 by Histidine (P23H mutant opsin protein) before said administering.